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NEWS 20 Feb 13 CANCERLIT is no longer being updated
NEWS 21 Feb 24 METADEX enhancements
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NEWS 29 Mar 24 Additional information for trade-named substances without
structures available in REGISTRY
NEWS 30 Apr 11 Display formats in DGENE enhanced
NEWS 31 Apr 14 MEDLINE Reload
NEWS 32 Apr 17 Polymer searching in REGISTRY enhanced
NEWS 33 Jun 13 Indexing from 1947 to 1956 added to records in CA/CAPLUS
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added to PHAR
NEWS 37 May 15 MEDLINE file segment of TOXCENTER reloaded
NEWS 38 May 15 Supporter information for ENCOMPPAT and ENCOMPLIT updated
NEWS 39 May 16 CHEMREACT will be removed from STN

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NEWS 40 May 19 Simultaneous left and right truncation added to WSCA
NEWS 41 May 19 RAPRA enhanced with new search field, simultaneous left and right truncation
NEWS 42 Jun 06 Simultaneous left and right truncation added to CBNB
NEWS 43 Jun 06 PASCAL enhanced with additional data
NEWS 44 Jun 20 2003 edition of the FSTA Thesaurus is now available

NEWS EXPRESS April 4 CURRENT WINDOWS VERSION IS V6.01a, CURRENT MACINTOSH VERSION IS V6.0b(ENG) AND V6.0Jb(JP), AND CURRENT DISCOVER FILE IS DATED 01 APRIL 2003

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FILE COVERS 1907 - 20 Jun 2003 VOL 138 ISS 26

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FILE LAST UPDATED: 19 Jun 2003 (20030619/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 17-methylene steroids
587388 17
108691 METHYLENE
103724 STEROIDS
L1 7 17-METHYLENE STEROIDS
(17(W)METHYLENE(W)STEROIDS)

=> s 11 full
587388 17
108691 METHYLENE
103724 STEROIDS
L2 7 17-METHYLENE STEROIDS
(17(W)METHYLENE(W)STEROIDS)

=> s 17-methylene steroids full
587388 17
108691 METHYLENE
103724 STEROIDS
L3 7 17-METHYLENE STEROIDS
(17(W)METHYLENE(W)STEROIDS)

=> d 12 1-7 ibib hitstr abs

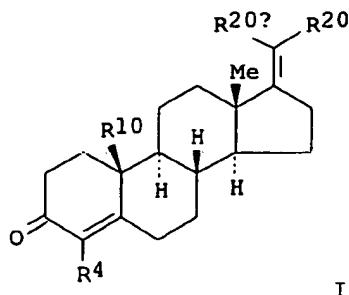
L2 ANSWER 1 OF 7 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 2002:184859 CAPLUS
DOCUMENT NUMBER: 136:247741
TITLE: Method for the production of 17-methylene steroids and pharmaceutical compositions containing them
INVENTOR(S): Menzenbach, Bernd; Elger, Walter; Droscher, Peter; Hillisch, Alexander; Kaufmann, Guenter; Schweikert, Hans-Udo; Mueller, Gerd
PATENT ASSIGNEE(S): Jenapharm G.m.b.H. & Co. K.-G., Germany
SOURCE: PCT Int. Appl., 28 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: German
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---------------|------|----------|-----------------|----------|
| WO 2002019971 | A1 | 20020314 | WO 2001-EP9943 | 20010829 |
| WO 2002019971 | C2 | 20020808 | | |

W: AE, AG, AU, BA, BB, BG, BR, CA, CN, CO, CR, CU, CZ, DM, DZ, EC, EE, GD, GE, HR, HU, ID, IL, IN, IS, JP, KP, KR, LC, LK, LR, LS, MA, MG, MN, MX, NO, NZ, PL, SG, SK, TT, UA, US, UZ, VN, YU, ZA, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

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| | | | | |
|------------------------|----|--|--------------------|----------|
| DE 10043846 | A1 | 20020404 | DE 2000-10043846 | 20000904 |
| AU 2002010470 | A5 | 20020322 | AU 2002-10470 | 20010829 |
| US 2002091112 | A1 | 20020711 | US 2002-963680 | 20020125 |
| NO 2003000989 | A | 20030502 | NO 2003-989 | 20030303 |
| PRIORITY APPLN. INFO.: | | | DE 2000-10043846 A | 20000904 |
| | | | US 2000-243281P P | 20001026 |
| | | | WO 2001-EP9943 W | 20010829 |
| OTHER SOURCE(S): | | CASREACT 136:247741; MARPAT 136:247741 | | |
| GI | | | | |



AB The inventive compds., e.g. I [R4 = halogen, pseudohalogen (CN, N3); R10 = H, straight or branched Cl-4-alkyl; R20, R20a = H, straight or branched Cl-4-alkyl, hydroxy-Cl-4-alkyl or one of R20, R20a = H, straight or branched Cl-4-alkyl, hydroxy-Cl-4-alkyl and the other is a halogen, pseudohalogen], have an active profile with a hybrid character of such that they act as inhibitors of the 5.alpha.-reductase and, at the same time, as gestagens. Thus, I (R4= R20 = Cl, R10 = H, R20a = H) was prep'd. from 17.alpha.- (chloromethyl)-17-hydroxyestr-4-en-3-one via dehydration with SOCl₂ in pyridine, regioselective epoxidn. and chlorination/dehydration. Said compds. are thus suited for treating medical disorders that, in men and women, are a result of an increased androgen level in certain organs and tissues. The inventive compds. combined with other hormonal substances such as estrogen, testosterone or a potent androgen are suited as contraceptives for women and men. Thus, I (R4= R20 = Cl, R10 = H, R20a = H) showed IC₅₀ = 250 nM vs. 5.alpha.-reductase.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1990:459681 CAPLUS
DOCUMENT NUMBER: 113:59681
TITLE: Steroidal cyclobutanones. I. The synthesis and stereochemistry of steroidal spirocyclobutanones
AUTHOR(S): Paryzek, Zdzislaw; Blaszczuk, Krzysztof
CORPORATE SOURCE: Fac. Chem., Adam Mickiewicz Univ., Poznan, 60-780, Pol.
SOURCE: Liebigs Annalen der Chemie (1990), (7), 665-70
DOCUMENT TYPE: CODEN: LACHDL; ISSN: 0170-2041
LANGUAGE: English
OTHER SOURCE(S): CASREACT 113:59681

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GI For diagram(s), see printed CA Issue.
AB Cycloaddn. of C12C:CO to 3-, 7-, and 17-methylene steroids gave spirodichlorocyclobutanones, which were reduced to monochloro- and unsubstituted spirocyclobutanones. Selective cycloaddn. to the exo-double bond was obsd. in the reaction of 3.beta.-acetoxy-17-methylene-5-androstene giving cyclobutanone I (R = Cl) which was reduced to I (R = H). H2O2 oxidn. of I (R = H) gave the lactone II. The stereochem. of the spiro compds. was assigned on the basis of 1H- and 13C-NMR spectra.

L2 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 1990:56425 CAPLUS
DOCUMENT NUMBER: 112:56425
TITLE: Preparation of 9.alpha.-hydroxy-17-methylene steroids as intermediates for corticosteroids
INVENTOR(S): Batist, Jacobus Nicolaas Maria; Marx, Arthur Friedrich
PATENT ASSIGNEE(S): Gist-Brocades N. V., Neth.
SOURCE: Eur. Pat. Appl., 45 pp.
CODEN: EPXXDW
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|---|------|----------|-----------------|----------|
| EP 336521 | A1 | 19891011 | EP 1989-200891 | 19890407 |
| EP 336521 | B1 | 19920401 | | |
| R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE | | | | |
| WO 8909781 | A1 | 19891019 | WO 1989-NL20 | 19890407 |
| W: AU, DK, FI, HU, JP, KR, NO, US | | | | |
| AU 8934313 | A1 | 19891103 | AU 1989-34313 | 19890407 |
| AU 618350 | B2 | 19911219 | | |
| HU 55411 | A2 | 19910528 | HU 1989-2602 | 19890407 |
| HU 208437 | B | 19931028 | | |
| JP 03503645 | T2 | 19910815 | JP 1989-504593 | 19890407 |
| AT 74363 | E | 19920415 | AT 1989-200891 | 19890407 |
| ES 2033516 | T3 | 19930316 | ES 1989-200891 | 19890407 |
| IL 89880 | A1 | 19940624 | IL 1989-89880 | 19890407 |
| CN 1036774 | A | 19891101 | CN 1989-102092 | 19890408 |
| CN 1032211 | B | 19960703 | | |
| CA 1332409 | A1 | 19941011 | CA 1989-596257 | 19890410 |
| NO 8904898 | A | 19891206 | NO 1989-4898 | 19891206 |
| DK 9002408 | A | 19901005 | DK 1990-2408 | 19901005 |
| NO 9004333 | A | 19901203 | NO 1990-4333 | 19901005 |
| US 5194602 | A | 19930316 | US 1990-474852 | 19901212 |
| CN 1141301 | A | 19970129 | CN 1995-120256 | 19951124 |
| PRIORITY APPLN. INFO.: | | | EP 1988-200675 | 19880408 |
| | | | EP 1989-200891 | 19890407 |
| | | | WO 1989-NL20 | 19890407 |

OTHER SOURCE(S): MARPAT 112:56425

GI For diagram(s), see printed CA Issue.

AB The title compds. [I; R1 = H, halo, cyano, isocyano, HCONH, alkoxy; R2 = NO₂, Me, alkoxy carbonyl, hydroxymethyl, alkylcarbonyloxymethyl; R3 = H; R4 = H, OH, Me; or R3R4 = CH₂; the steroid nucleus may contain double bonds and further substituents; exception being 9.alpha.,21-dihydroxypregna-

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4,17(20)-diene-3,11-dione and its 21-acetate], useful as intermediate for corticosteroid, are prep'd. 3,3-(Ethylenedioxy)-9. α .-hydroxyandrost-5-en-17-one was refluxed with MeNO₂ in H₂NCH₂CH₂NH₂ for 24 h to give 3,3-(ethylenedioxy)-17-(nitromethylene)androst-5-en-9. α -ol.

L2 ANSWER 4 OF 7 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1981:462492 CAPLUS

DOCUMENT NUMBER: 95:62492

TITLE: D-Homo steroids from oxidation of 17-methylene steroids by thallium(III) nitrate

AUTHOR(S): Forcellese, Maria Luigia; Camerini, Elio; Ruffini, Bruna; Mincione, Enrico

CORPORATE SOURCE: Cent. Stud. Chim. Sostanze Org. Nat., CNR, Italy

SOURCE: Journal of Organic Chemistry (1981), 46(16), 3326-8

CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Thallium (III)-nitrate reacts with 17-methylene steroids to form D-homo-17. α .-methoxy-17a-oxo compds. via ring enlargement, enolization, oxythallation, and methanolysis.

L2 ANSWER 5 OF 7 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1977:140337 CAPLUS

DOCUMENT NUMBER: 86:140337

TITLE: Single and triple Vilsmeier formylation of 17-methylene steroids

AUTHOR(S): Dauphin, G.; Planat, D.

CORPORATE SOURCE: Equipe Rech. Assoc. CNRS No. 392, Univ. Clermont, Aubiere, Fr.

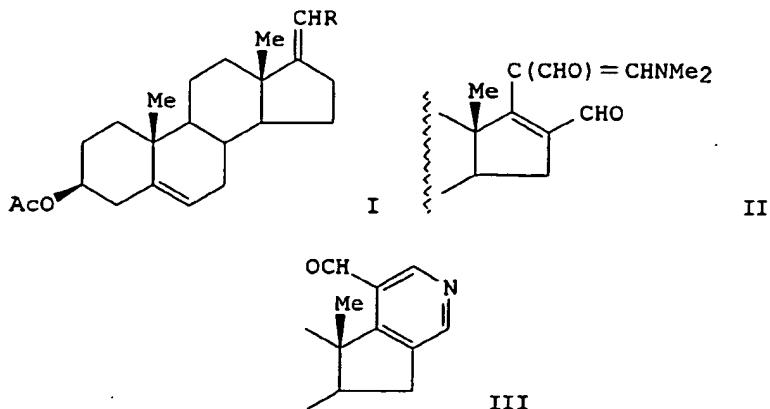
SOURCE: Tetrahedron Letters (1976), (45), 4065-8

CODEN: TELEAY; ISSN: 0040-4039

DOCUMENT TYPE: Journal

LANGUAGE: French

GI



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AB The androstene I (R = H) with DMF-POCl₃ in the ratios 1:15 and 1:1.5 for 15 and 1 day, resp. gave 50% 20 β -pregnatriene II and 40% 17(20)-E-pregnadiene I (R = CHO), resp. Analogous products were derived from the 5. α -H-5,6-dihydro analog of I(R = H). II and its 5,6-dihydro analog with ethanolic NH₃ gave 80-90% pyridoandrostanes III.

L2 ANSWER 6 OF 7 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1976:165101 CAPLUS
DOCUMENT NUMBER: 84:165101
TITLE: Oxidation of 17-methylene steroids by thallium(III) and mercury(II) acetates
AUTHOR(S): Ortar, G.; Arpiani, M. P.; Romeo, A.
CORPORATE SOURCE: Cent. Stud. Chim. Farm., Cons. Naz. Ric., Rome, Italy
SOURCE: Steroids (1976), 27(2), 197-203
CODEN: STEDAM; ISSN: 0039-128X
DOCUMENT TYPE: Journal
LANGUAGE: English

AB The reaction of 17-methyleneandrostanes with Tl(OAc)₃ in hot AcOH resulted in the formation of a mixt. of allylic compds. Oxymercuration in Me₃COH followed by reductive demercuration gave 17-methylene-16. β -hydroxyandrostanes as the major products.

L2 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1961:144432 CAPLUS
DOCUMENT NUMBER: 55:144432
ORIGINAL REFERENCE NO.: 55:27433f-h
TITLE: 16. α -Monohalomethyl steroids
INVENTOR(S): Kaspar, Emanuel; Wiechert, Rudolf; Schenck, Martin
PATENT ASSIGNEE(S): Schering Akt.-Ges
DOCUMENT TYPE: Patent
LANGUAGE: Unavailable
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

| PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------------|------|----------|-----------------|------|
| DE 1096903 | | 19610112 | DE | |
| GB 937616 | | | GB | |
| US 3232961 | | 1966 | US | |

AB 16,17-Methylene steroids of the 20-oxopregnane series were treated with a hydrohalide to give the title compds., useful as pharmaceuticals and intermediates in the manuf. of such. Thus, 200 mg. 16,17-methylene-5. α -pregnan-3. β -ol-20-one in 20 cc. CH₂Cl₂ was satd. with gaseous HBr, the mixt. kept 30 min. at room temp., washed, dried, and concd. to give 16. α -bromomethyl-5. α -pregnan-3. β -ol-20-one, m. 169 70.degree. (iso-Pr₂O), [α .D]20D 55.degree.. Similarly were prep'd.: 16. α -chloromethyl-5. α -pregnan-3. β -ol-20-one, m. 174-5.degree. (hexane), [α .D]28D 58.degree.; 16. α -iodomethyl-5. α -pregnan-3. β -ol-20-one [3-formate m. 138-9.degree. (iso-Pr₂O), [α .D]25D 32.5.degree.]; 16. α -iodomethyl-5-pregn-3. β -ol-20-one [3-formate m. 141-4.degree. (MeOH), [α .D]26D 9.1.degree.]; 16. α -chloromethyl-5-pregn-3. β -ol-20-one, m. 184-6.5.degree. (iso-Pr₂O).

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ENTRY | TOTAL
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